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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/534,314	10/24/2005	Anthony Rosenzweig	00786/431002 9779 EXAMINER	
21559	7590 05/25/2006			
CLARK & ELBING LLP			WHITEMAN, BRIAN A	
101 FEDERAL STREET BOSTON, MA 02110			ART UNIT	PAPER NUMBER
ŕ			1635	
			DATE MAILED: 05/25/2006	

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)			
	10/534,314	ROSENZWEIG ET AL.			
Office Action Summary	Examiner	Art Unit			
	Brian Whiteman	1635			
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address			
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).  Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).					
Status					
1) Responsive to communication(s) filed on  2a) This action is <b>FINAL</b> . 2b) This  3) Since this application is in condition for allowan closed in accordance with the practice under <i>E</i>	action is non-final. ace except for formal matters, pro				
Disposition of Claims					
4) ⊠ Claim(s) 15,16,20,21,29,30,42,45-49,51,54 and 4a) Of the above claim(s) is/are withdraw 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) 15,16,20,21,29,30,42,45-49,51,54,55	vn from consideration.				
Application Papers					
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) access Applicant may not request that any objection to the confidence of Replacement drawing sheet(s) including the correction 11) The oath or declaration is objected to by the Examine 11.	epted or b) objected to by the Eddrawing(s) be held in abeyance. See ion is required if the drawing(s) is obj	e 37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).  a) All b) Some * c) None of:  1. Certified copies of the priority documents have been received.  2. Certified copies of the priority documents have been received in Application No  3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).  * See the attached detailed Office action for a list of the certified copies not received.					
Attachment(s)  1) Notice of References Cited (PTO-892)  2) Notice of Draftsperson's Patent Drawing Review (PTO-948)  3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date	4)  Interview Summary Paper No(s)/Mail Da 5)  Notice of Informal P 6)  Other:				

## **DETAILED ACTION**

Claims 15, 16, 20, 21, 29, 30, 42, 45-49, 51, 54, and 55 are pending.

The cancellation of claims 1-14, 17-19, 22-28, 31-41, 43, 44, 50, 52, 53 and 56-64, the amendment to claims 16, 20, 21, 46-48, and 55 in paper filed on 5/9/05 is acknowledged.

## Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 15, 16, and 21, drawn to a method of treating a cardiac disorder in a mammal comprising administering to said mammal a FADD protein.

Group II, claim(s) 15, 16, 20 and 21, drawn to a method of treating a cardiac disorder in a mammal comprising administering to said mammal an anti-inflammatory FADD inhibitor.

Group III, claim(s) 29, drawn to a human cardiomyocyte expressing a dominant negative FADD protein.

Group IV, claim(s) 30, drawn to a human cardiomyocyte expressing a recombinant FADD protein.

Group V, claim(s) 42, 45, 46, 47, 48, and 49, drawn to a method of identifying a candidate compound for treating cardiac inflammation in a mammal comprising contacting a

Art Unit: 1635

cardiomyocyte expressing a FADD gene candidate compound and measuring FADD gene expression in said cardiomyocyte.

Group VI, claim(s) 42, 45, 46, 47, 48, and 49, drawn to a method of identifying a candidate compound for treating cardiac inflammation in a mammal comprising contacting a cardiomyocyte expressing a FADD gene candidate compound and measuring FADD protein activity in said cardiomyocyte.

Group VII, claim(s) 51, 54, and 55, drawn to a method for identifying a candidate compound for treating cardiac inflammation or cardiac disorder comprising contacting a FADD protein with a candidate compound and determining whether compound bind said FADD protein.

The inventions listed as Groups I-VII do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical feature for the following reasons:

37 CFR 1.475(b) states:

"An international or a national stage application containing claims to different categories of invention will be considered to have unity of invention if the claims are drawn only to one of the following combinations of categories:

- (1) A product and a process specially adapted for the manufacture of said product; or
- (2) A product and process of use of said product; or
- (3) A product, a process specially adapted for the manufacture of the said product, and a use of the said product; or

Application/Control Number: 10/534,314 Page 4

Art Unit: 1635

(4) A process and an apparatus or means specifically designed for carrying out the said process; or

(5) A product, a process specially adapted for the manufacture of the said product, and an apparatus or means specifically designed for carrying out the said process.

37 CFR 1.475(c) states:

"If an application contains claims to more or less than one of the combination of categories of invention set forth in paragraph (b) of this section, unity of invention might not be present."

37 CFR 1.475(d) also states:

"If multiple products, processes of manufacture or uses are claimed, the first invention of the category first mentioned in the claims of the application and the first recited invention of each other categories related thereto will be considered as the main invention in the claims, see PCT Article 17(3)(a) and 1.476(c)."

37 CFR 1.475(e) further states:

"The determination whether a group of inventions is so linked as to form a single general inventive concept shall be made without regard to whether the inventions are claimed in separate claims or as alternative within a single claim."

In view of 37 CFR 1.475 (b), 37 CFR 1.475 (c), 37 CFR 1.475 (d), and 37 CFR 1.475 (e), Group I is considered the main invention to the product first mentioned in the claims, and the first recited invention drawn to other categories related thereto, e.g. a method of making, method of use.

Application/Control Number: 10/534,314

Art Unit: 1635

The technical feature linking groups I-VII appears to be a cardiomyocyte expressing a recombinant FADD protein.

However, Chao et al. (JBC, 277:31639-31645, 2002) teaches a cardiomyocyte comprising a wild type or dominant negative form of FADD.

Therefore, the technical feature linking the inventions of groups I-VII does not constitute a special technical feature as defined by PCT Rule 13.2, as it does not define a contribution over the prior art.

The special technical feature of Group I is considered to be a method of treating a cardiac disorder in a mammal comprising administering to said mammal a FADD protein.

The special technical feature of Group II is considered to be a method of treating a cardiac disorder in a mammal comprising administering to said mammal an anti-inflammatory FADD inhibitor.

The special technical feature of Group III is considered to be a human cardiomyocyte expressing a dominant negative FADD protein.

The special technical feature of Group IV is considered to be a human cardiomyocyte expressing a recombinant FADD protein.

The special technical feature of Group V is considered to be a method of identifying a candidate compound for treating cardiac inflammation in a mammal comprising contacting a cardiomyocyte expressing a FADD gene candidate compound and measuring FADD gene expression in said cardiomyocyte.

The special technical feature of Group VI is considered to be to a method of identifying a candidate compound for treating cardiac inflammation in a mammal comprising contacting a

cardiomyocyte expressing a FADD gene candidate compound and measuring FADD protein activity in said cardiomyocyte.

Page 6

The special technical feature of Group VII is considered to be a method for identifying a candidate compound for treating cardiac inflammation or cardiac disorder comprising contacting a FADD protein with a candidate compound and determining whether compound bind said FADD protein.

Accordingly, Groups I-VII are not so linked by the same or a corresponding technical feature as to form a single general inventive concept.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Brian Whiteman whose telephone number is (571) 272-0764. The examiner can normally be reached on Monday through Friday from 7:00 to 4:00 (Eastern Standard Time), with alternating Fridays off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Peter Paras, SPE – Art Unit 1635, can be reached at (571) 272-4517.

Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Group 1600 via the PTO Fax Center. The faxing of

Application/Control Number: 10/534,314 Page 7

Art Unit: 1635

such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989). The Fax Center number is (571) 273-8300.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to (571) 272-0547.

Patent applicants with problems or questions regarding electronic images that can be viewed in the Patent Application Information Retrieval system (PAIR) can now contact the USPTO's Patent Electronic Business Center (Patent EBC) for assistance. Representatives are available to answer your questions daily from 6 am to midnight (EST). The toll free number is (866) 217-9197. When calling please have your application serial or patent number, the type of document you are having an image problem with, the number of pages and the specific nature of the problem. The Patent Electronic Business Center will notify applicants of the resolution of the problem within 5-7 business days. Applicants can also check PAIR to confirm that the problem has been corrected. The USPTO's Patent Electronic Business Center is a complete service center supporting all patent business on the Internet. The USPTO's PAIR system provides Internet-based access to patent application status and history information. It also enables applicants to view the scanned images of their own application file folder(s) as well as general patent information available to the public.

For all other customer support, please call the USPTO Call Center (UCC) at 800-786-9199.

Brian Whiteman

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